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(54) Polymer wood composite.

*we don't have this
we use PE*

(57) A polymer and wood fibre composite that can be used in the form of a linear extrudate or thermoplastic pellet to manufacture structural members, for example in an extrusion process or an injection moulding process. The composite comprises a vinyl chloride polymer, wood fibre and water, and preferably also intentionally recycled impurity which might include thermoplastic polymer, adhesive, paint, or thermoplastic resin. The pellet can have a cross-section shape having a volume of at least about 12 mm³. Preferably the pellet is a right cylindrical pellet having a minimum radius of about 1.5 mm and a minimum length of 1 mm weighing at least 14 mg.

This invention relates to composite thermoplastic materials used for the fabrication of structural members. The composite material can be provided in pellet form, for use in any structural unit; preferably they are for use in windows and doors for residential and commercial architecture.

These composite materials include an optional, intentional recycle of by-product streams, from window and door manufacture, of polymer, adhesive, paint, etc. More particularly, the invention relates to an improved composite material adapted to extrusion or injection moulding processes for forming structural members that have improved properties when used in windows and doors. The composite materials of the invention can be made to manufacture structural components such as rails, jambs, stiles, sills, tracks, stop and sash, nonstructural trim elements such as grid, cove, bead, quarter round, etc.

Conventional window and door manufacture has commonly used wood and metal components in forming structural members. Commonly, residential windows are manufactured from milled wood products that are assembled with glass to form typically double hung or casement units. Wood windows while structurally sound, useful and well adapted for use in many residential installations, can deteriorate under certain circumstances. Wood windows also require painting and other periodic maintenance. Wooden windows also suffer from cost problems related to the availability of suitable wood for construction. Clear wood products are slowly becoming more scarce and are becoming more expensive as demand increases. Metal components are often combined with glass and formed into single unit sliding windows. Metal windows typically suffer from substantial energy loss during winter months.

Extruded thermoplastic materials have been used in window and door manufacture. Filled and unfilled thermoplastics have been extruded into useful seals, trim, weatherstripping, coatings and other window construction components. Thermoplastic materials such as polyvinyl chloride have been combined with wood members in manufacturing windows sold by Andersen Corporation for many years under the trade mark PER-MASHIELD. The technology used in the production of such windows is disclosed in US-2926729 and US-3432883, and is used in the application plastic coatings or envelopes on wooden or other structural members. Generally, the cladding or coating technology used in making such windows involves extruding a thin polyvinyl chloride coating or envelope surrounding a wooden structural member.

Polyvinyl chloride has been combined with wood to make extruded materials. However, such materials have not successfully been used in the form of a structural member that is a direct replacement for wood. Common thermoplastic composite materials cannot provide similar thermal and structural properties to wood. These extruded materials fail to have sufficient modulus, compressive strength, coefficient of thermal expansion that matches wood to produce a direct replacement material. Further, many prior art extruded composites must be milled after extrusion to a final useful shape. Typical commodity plastics have achieved a modulus no greater than about 500,000 psi (3400 MPa). One class of composite, a polyvinyl chloride/wood flour material, poses the added problem that wood dust, which can accumulate during manufacture, tends to be explosive at certain concentrations of wood flour in the air. Most commonly, poly vinyl chloride, polystyrene and polyethylene thermoplastics have been used in such products.

Accordingly, a substantial need exists for a composite material that can be made of polymer and wood fibre with an optional, intentional recycle of a waste stream. A further need exists for a composite material that can be extruded into a shape that is a direct substitute for the equivalent milled shape in a wooden or metal structural member. This need requires a coefficient of thermal expansion that approximates wood, a material that can be extruded into reproducible stable dimensions, a high compressive strength, a low thermal transmission rate, an improved resistance to insect attack and rot while in use and a hardness and rigidity that permits sawing, milling, and fastening retention comparable to wood members. Further, companies manufacturing window and door products have become significantly sensitive to waste streams produced in the manufacture of such products. Substantial quantities of wood waste including wood trim pieces, sawdust, wood milling by-products; recycled thermoplastic including recycled polyvinyl chloride, has caused significant expense to window manufacturers. Commonly, these materials are either burned for their heat value in electrical generation or are shipped to qualified landfills for disposal. Such waste streams are contaminated with substantial proportions of hot melt and solvent-based adhesives, waste thermoplastic such as polyvinyl chloride, paint, preservatives, and other organic materials. A substantial need exists to find a productive environmentally compatible use for such waste streams to avoid returning the materials into the environment in an environmentally harmful way.

It has been found that the problems relating to forming a substitute for wood and metal structural members and the problems relating to the recycle of waste streams in window manufacture can be solved by forming a polymer/wood fibre composite material into window and door structural members. The polyvinyl chloride can be combined with wood fibre and waste materials to form a polyvinyl chloride/wood fibre composite, preferably in the form of a pellet. The wood fibre comprises the sawdust or milling byproduct waste stream from milling wooden members in window manufacture and can be contaminated with substantial proportions of hot melt

adhesive, paint, solvent or adhesive components, preservatives, polyvinyl chloride recycle pigment, plasticizers, etc. We have found that the PVC and wood fibre composite can be manufactured into acceptable substitutes for wooden members if the PVC and wood material contains less than about 10 wt %, preferably less than 3.5% water based on pellet weight. The compositions can achieve in a final product high modulus, high compressive strength, reproducible, stable dimensions and a superior modulus of elasticity. It has also been found that the successful manufacture of structural members for windows and doors requires the preliminary manufacture of the polyvinyl chloride wood fibre composite in the form of a pellet wherein the materials are intimately mixed and contacted in forming the pellet prior to the extrusion of the members from the pellet material. It has been found that the intimate mixing of the polyvinyl chloride, wood fibre, and waste in the manufacture of the pellet process with associated control of moisture content produces a pelletized product that is uniquely adapted to the extrusion manufacture of PVC/wood fibre components and achieves the manufacture of a useful wood replacement product.

The invention relates to the use of polyvinyl chloride and wood fibre composite materials with a controlled water content in the form of a pelletized material wherein the wood fibre is intimately contacted and wetted by the organic materials. The intimate contact and wetting between the components in the pelletizing process ensures high quality physical properties in the extruded composite materials after manufacture.

Pellet

The polyvinyl chloride and wood fibre can be combined and formed into a pellet using a thermoplastic extrusion processes. Wood fibre can be introduced into pellet making process in a number of sizes. We believe that the wood fibre should have a minimum size of length and width of at least 1 mm because wood flour tends to be explosive at certain wood to air ratios. Further, wood fibre of appropriate size of a aspect ratio greater than 1 tends to increase the physical properties of the extruded structural member. However, useful structural members can be made with a fibre of very large size. Fibres that are up to 3 cm in length and 0.5 cm in thickness can be used as input to the pellet or linear extrudate manufacturing process. However, particles of this size do not produce highest quality structural members or maximized structural strength. The best appearing product with maximized structural properties are manufactured within a range of particle size as set forth below. Further, large particle wood fibre can be reduced in size by grinding or other similar processes that produce a fibre similar to sawdust having the stated dimensions and aspect ratio. One further advantage of manufacturing sawdust of the desired size is that the material can be pre-dried before introduction into the pellet or linear extrudate manufacturing process. Further, the wood fibre can be pre-pelletized into pellets of wood fibre with small amounts of binder if necessary.

During the pelletizing process for the composite pellet, the polyvinyl chloride and wood fibre are intimately contacted at high temperatures and pressures to insure that the wood fibre and polymeric material are wetted, mixed and extruded in a form such that the polymer material, on a microscopic basis, coats and flows into the pores, cavity, etc., of the fibres. The fibres are preferably substantially oriented by the extrusion process in the extrusion direction. Such substantial orientation causes overlapping of adjacent parallel fibres and polymeric coating of the oriented fibres resulting a material useful for manufacture of improved structural members with improved physical properties. The degree of orientation is about 20%, preferably 30% above random orientation which is about 45 to 50%. The structural members have substantially increased strength and tensile modulus with a coefficient of thermal expansion and a modulus of elasticity that is optimized for window and doors. The properties are a useful compromise between wood, aluminum and neat polymer.

Moisture control is an important element of manufacturing a useful linear extrudate or pellet. Depending on the equipment used and processing conditions, control of the water content of the linear extrudate or pellet can be important in forming a successful structural member substantially free of internal voids or surface blemishes. The concentration of water present in the sawdust during the formation of pellet or linear extrudate when heated can flash from the surface of the newly extruded structural member and can come as a result of a rapid volatilization, form a steam bubble deep in the interior of the extruded member which can pass from the interior through the hot thermoplastic extrudate leaving a substantial flaw. In a similar fashion, surface water can bubble and leave cracks, bubbles or other surface flaws in the extruded member.

Trees when cut depending on relative humidity and season can contain from 30 to 300 wt % water based on fibre content. After rough cutting and finishing into sized lumber, seasoned wood can have a water content of from 20 to 30 wt % based on fibre content. Kiln dried sized lumber cut to length can have a water content typically in the range of 8 to 12%, commonly 8 to 10 wt % based on fibre. Some wood source, such as poplar or aspen, can have increased moisture content while some hard woods can have reduced water content.

Because of the variation in water content of wood fibre source and the sensitivity of extrudate to water content control of water to a level of less than 8 wt % in the pellet based on pellet weight is important. Structural

members extruded in non-vented extrusion process, the pellet should be as dry as possible and have a water content between 0.01 and 5%, preferably less than 3.5 wt %. When using vented equipment in manufacturing the extruded linear member, a water content of less than 8 wt % can be tolerated if processing conditions are such that vented extrusion equipment can dry the thermoplastic material prior to the final formation of the structural member of the extrusion head.

The pellets or linear extrudate of the invention are made by extrusion of the polyvinyl chloride and wood fibre composite through an extrusion die resulting in a linear extrudate that can be cut into a pellet shape. The pellet cross-section can be any arbitrary shape depending on the extrusion die geometry. However, we have found that a regular geometric cross-sectional shape can be useful. Such regular cross-sectional shapes include a triangle, a square, a rectangle, a hexagonal, an oval, a circle, etc. The preferred shape of the pellet is a regular cylinder having a roughly circular or somewhat oval cross-section. The pellet volume is preferably greater than about 12 mm³. The preferred pellet is a right circular cylinder, the preferred radius of the cylinder is at least 1.5 mm with a length of at least 1 mm. Preferably, the pellet has a radius of 1 to 5 mm and a length of 1 to 10 mm. Most preferably, the cylinder has a radius of 2.3 to 2.6 mm, a length of 2.4 to 4.7 mm, a volume of 40 to 100 mm³, a weight of 40 to 130 mg and a bulk density of about 0.2 to 0.8 g.mm⁻³.

It has been found that the interaction, on a microscopic level, between the polymer mass and the wood fibre is an important element of the invention. We have found that the physical properties of an extruded member are improved when the polymer melt during extrusion of the pellet or linear member thoroughly wets and penetrates the wood fibre particles. The thermoplastic material comprises an exterior continuous organic polymer phase with the wood particle dispersed as a discontinuous phase in the continuous polymer phase. The material during mixing and extrusion obtains an aspect ratio of at least 1.1 and preferably between 2 and 4, optimizes orientation such as at least 20 wt %, preferably 30% of the fibres are oriented in an extruder direction and are thoroughly mixed and wetted by the polymer such that all exterior surfaces of the wood fibre are in contact with the polymer material. This means, that any pore, crevice, crack, passage way, indentation, etc., is fully filled by thermoplastic material. Such penetration is attained by ensuring that the viscosity of the polymer melt is reduced by operations at elevated temperature and the use of sufficient pressure to force the polymer into the available internal pores, cracks and crevices in and on the surface of the wood fibre.

During the pellet or linear extrudate manufacture, substantial work is done in providing a uniform dispersion of the wood into the polymer material. Such work produces substantial orientation which when extruded into a final structural member, permits the orientation of the fibres in the structural member to be increased in the extruder direction resulting in improved structural properties.

The pellet dimensions are selected for both convenience in manufacturing and in optimizing the final properties of the extruded materials. A pellet is with dimensions substantially less than the dimensions set forth above are difficult to extrude, pelletize and handle in storage. Pellets larger than the range recited are difficult to introduce into extrusion or injection moulding equipment, and are difficult to melt and form into a finished structural member.

Polyvinyl chloride homopolymer, copolymers and polymeric alloys

Polyvinyl chloride is a common commodity thermoplastic polymer. Vinyl chloride monomer is made from a variety of different processes such as the reaction of acetylene and hydrogen chloride and the direct chlorination of ethylene. Polyvinyl chloride is typically manufactured by the free radical polymerization of vinyl chloride resulting in a useful thermoplastic polymer. After polymerization, polyvinyl chloride is commonly combined with thermal stabilizers, lubricants, plasticizers, organic and inorganic pigments, fillers, biocides, processing aids, flame retardants and other commonly available additive materials. Polyvinyl chloride can also be combined with other vinyl monomers in the manufacture of polyvinyl chloride copolymers. Such copolymers can be linear copolymers, branched copolymers, graft copolymers, random copolymers, regular repeating copolymers, block copolymers, etc. Monomers that can be combined with vinyl chloride to form vinyl chloride copolymers include acrylonitrile; alpha-olefins such as ethylene, propylene, etc.; chlorinated monomers such as vinylidene dichloride, acrylate monomers such as acrylic acid, methylacrylate, methylmethacrylate, acrylamide, hydroxyethyl acrylate, and others; styrenic monomers such as styrene, alphamethyl styrene, vinyl toluene, etc.; vinyl acetate; and other commonly available ethylenically unsaturated monomer compositions.

Such monomers can be used in an amount of up to about 50 mol %, the balance being vinyl chloride. Polymer blends or polymer alloys can be useful in manufacturing the pellet or linear extrudate of the invention. Such alloys typically comprise two miscible polymers blended to form a uniform composition. Scientific and commercial progress in the area of polymer blends has lead to the realization that important physical property improvements can be made not by developing new polymer material but by forming miscible polymer blends or alloys. A polymer alloy at equilibrium comprises a mixture of two amorphous polymers existing as a single phase

of intimately mixed segments of the two macro molecular components. Miscible amorphous polymers form glasses upon sufficient cooling and a homogeneous or miscible polymer blend exhibits a single, composition dependent glass transition temperature (T_g). Immiscible or non-alloyed blend of polymers typically displays two or more glass transition temperatures associated with immiscible polymer phases. In the simplest cases, the properties of polymer alloys reflect a composition weighted average of properties possessed by the components. In general, however, the property dependence on composition varies in a complex way with a particular property, the nature of the components (glassy, rubbery or semi-crystalline), the thermodynamic state of the blend, and its mechanical state whether molecules and phases are oriented. Polyvinyl chloride forms a number of known polymer alloys including, for example, polyvinyl chloride/nitrile rubber; polyvinyl chloride and related chlorinated copolymers and terpolymers of polyvinyl chloride or vinylidene dichloride; polyvinyl chloride/alpha-methyl styrene-acrylonitrile copolymer blends; polyvinyl chloride/polyethylene; polyvinyl chloride/chlorinated polyethylene and others.

The primary requirement for the substantially thermoplastic polymeric material is that it retain sufficient thermoplastic properties to permit melt blending with wood fibre, permit formation of linear extrudate pellets, and to permit the composition material or pellet to be extruded or injection moulded in a thermoplastic process forming the rigid structural member. Polyvinyl chloride homopolymers copolymers and polymer alloys are available from a number of manufacturers including B.F. Goodrich, Vista, Air Products, Occidental Chemicals, etc. Preferred polyvinyl chloride materials are polyvinyl chloride homopolymer having a molecular weight of about $90,000 \pm 50,000$, most preferably about $88,000 \pm 10,000$.

Wood fibre

Wood fibre, in terms of abundance and suitability can be derived from either soft woods or evergreens or from hard woods commonly known as broad leaf deciduous trees. Soft woods are generally preferred for fibre manufacture because the resulting fibres are longer, contain high percentages of lignin and lower percentages of hemicellulose than hard woods. While soft wood is the primary source of fibre for the invention, additional fibre make-up can be derived from a number of secondary or fibre reclaim sources including bamboo, rice, sugar cane, and recycled fibres from newspapers, boxes, computer printouts, etc.

However, the primary source for wood fibre of this invention comprises the wood fibre by-product of sawing or milling soft woods commonly known as sawdust or milling tailings. Such wood fibre has a regular reproducible shape and aspect ratio. The fibres based on a random selection of about 100 fibres are commonly at least 3 mm in length, 1 mm in thickness and commonly have an aspect ratio of at least 1.8. Preferably, the fibres are 1 to 10 mm in length, 0.3 to 1.5 mm in thickness with an aspect ratio between 2 and 7, preferably 2.5 to 6.0. The preferred fibre for use in this invention are fibres derived from processes common in the manufacture of windows and doors. Wooden members are commonly ripped or sawed to size in a cross grain direction to form appropriate lengths and widths of wood materials. The by-product of such sawing operations is a substantial quantity of sawdust. In shaping a regular shaped piece of wood into a useful milled shape, wood is commonly passed through machines which selectively removes wood from the piece leaving the useful shape. Such milling operations produces substantial quantities of sawdust or mill tailing by-products. Lastly, when shaped materials are cut to size and mitred joints, butt joints, overlapping joints, mortise and tenon joints are manufactured from pre-shaped wooden members, substantial waste trim is produced. Such large trim pieces are commonly cut and machined to convert the larger objects into wood fibre having dimensions approximating sawdust or mill tailing dimensions. The wood fibre sources of the invention can be blended regardless of particle size and used to make the composite. The fibre stream can be pre-sized to a preferred range or can be sized after blending. Further, the fibre can be pre-pelletized before use in composite manufacture.

Such sawdust material can contain substantial proportions of waste stream by-products. Such by-products include waste polyvinyl chloride or other polymer materials that have been used as coating, cladding or envelope on wooden members; recycled structural members made from thermoplastic materials; polymeric materials from coatings; adhesive components in the form of hot melt adhesives, solvent based adhesives, powdered adhesives, etc.; paints including water based paints, alkyd paints, epoxy paints, etc.; preservatives, anti-fungal agents, anti-bacterial agents, insecticides, etc., and other waste streams common in the manufacture of wooden doors and windows. The total waste stream content of the wood fibre materials is commonly less than 25 wt % of the total wood fibre input into the polyvinyl chloride wood fibre product. Of the total waste recycle, approximately 10 wt % of that can comprise a vinyl polymer commonly polyvinyl chloride. Commonly, the intentional recycle ranges from about 1 to about 25 wt %, preferably about 2 to about 20 wt %, most commonly from about 3 to about 15 wt % of contaminants based on the sawdust.

Composition and pellet manufacture

In the manufacture of the composition and pellet of the invention, the manufacture and procedure requires two important steps. A first blending step and a second pelletizing step.

5 During the blending step, the polymer and wood fibre are intimately mixed by high shear mixing components with recycled material to form a polymer wood composite wherein the polymer mixture comprises a continuous organic phase and the wood fibre with the recycled materials forms a discontinuous phase suspended or dispersed throughout the polymer phase. The manufacture of the dispersed fibre phase within a continuous polymer phase requires substantial mechanical input. Such input can be achieved using a variety of mixing means including preferably extruder mechanisms wherein the materials are mixed under conditions of high shear until the appropriate degree of wetting and intimate contact is achieved. After the materials are fully mixed, the moisture content can be controlled at a moisture removal station. The heated composite is exposed to atmospheric pressure or reduced pressure at elevated temperature for a sufficient period of time to remove moisture resulting in a final moisture content of about 8 wt % or less. Lastly, the polymer fibre is aligned and extruded into a useful form.

10 The preferred equipment for mixing and extruding the composition and wood pellet of the invention is an industrial extruder device. Such extruders can be obtained from a variety of manufacturers including Cincinnati Millicron, etc.

15 The materials feed to the extruder can comprise from about 30 to 50 wt % of sawdust including recycled impurity along with from about 50 to 70 wt % of polyvinyl chloride polymer compositions. Preferably, about 35 to 45 wt % wood fibre or sawdust is combined with 65 to 55 wt % polyvinyl chloride homopolymer. The polyvinyl chloride feed is commonly in a small particulate size which can take the form of flake, pellet, powder, etc. Any polymer form can be used such that the polymer can be dry mixed with the sawdust to result in a substantially uniform pre-mix. The wood fibre or sawdust input can be derived from a number of plant locations including the sawdust resulting from rip or cross grain sawing, milling of wood products or the intentional commuting or fibre manufacture from waste wood scrap. Such materials can be used directly from the operations resulting in the wood fibre by-product or the by-products can be blended to form a blended product. Further, any wood fibre material alone, or in combination with other wood fibre materials, can be blended with waste stream by-product from the manufacturer of wood windows as discussed above. The wood fibre or sawdust can be combined with other fibres and recycled in commonly available particulate handling equipment.

20 Polymer and wood fibre are then dry blended in appropriate proportions prior to introduction into blending equipment. Such blending steps can occur in separate powder handling equipment or the polymer fibre streams can be simultaneously introduced into the mixing station at appropriate feed ratios to ensure appropriate product composition.

25 In a preferred mode, the wood fibre is placed in a hopper, controlled by weight or by volume, to meter the sawdust at a desired volume while the polymer is introduced into a similar hopper have a volumetric metering input system. The volumes are adjusted to ensure that the composite material contains appropriate proportions on a weight basis of polymer and wood fibre. The fibres are introduced into a twin screw extrusion device. The extrusion device has a mixing section, a transport section and melt section. Each section has a desired heat profile resulting in a useful product. The materials are introduced into the extruder at a rate of about 600 to about 1000 pounds of material per hour and are initially heated to a temperature of about 215-225°C. In the intake section, the stage is maintained at about 215°C to 225°C. In the mixing section, the temperature of the twin screw mixing stage is staged beginning at a temperature of about 205 to 215°C leading to a final temperature in the melt section of about 195 to 205°C at spaced stages. Once the material leaves the blending stage, it is introduced into a three stage extruder with a temperature in the initial section of 185 to 195°C wherein the mixed thermoplastic stream is divided into a number of cylindrical streams through a head section and extruded in a final zone of 195 to 200°C. Such head sections can contain a circular distribution (6 to 8" diameter) of 10 to 500 or more, preferably 20 to 250 orifices having a cross-sectional shape leading to the production of a regular cylindrical pellet. As the material is extruded from the head it is cut with a double-ended knife blade at a rotational speed of about 100 to 400 rpm resulting in the desired pellet length.

30 The following examples were performed to further illustrate the invention that is explained in detail above. The following information illustrates the typical production conditions and compositions and the tensile modulus of a structural member made from the pellet.

35 **EXAMPLE 1**

A Cincinnati millicron extruder with an HP barrel, Cincinnati pelletizer screws, an AEG K-20 pelletizing head with 260 holes, each hole having a diameter of about 0.0200 inches was used to make the pellet. The input to

the pelletizer comprised approximately 60 wt % polymer and 40 wt % sawdust. The polymer material comprises a thermoplastic mixture of approximately 100 parts of polyvinyl chloride homopolymer (in. weight of 88,000 ± 2000), about 15 parts titanium dioxide, about 2 parts ethylene bis-stearamide wax lubricant, about 1.5 parts calcium stearate, about 7.5 parts Rohm & Haas 820-T acrylic resin impact modifier/process aid and about 2 parts of dimethyl tin thioglycolate. The sawdust comprises a wood fibre particle containing about 5 wt % recycled polyvinyl chloride having a composition substantially identical to that recited above.

5 The initial melt temperature in the extruder was maintained between 350°C and 400°C. The pelletizer was operated at a polar vinyl chloride-sawdust composite combined through put of 800 pounds per hour. In the initial extruder feed zone, the barrel temperature was maintained between 215-225°C. In the intake zone, the barrel
10 was maintained at 215-225°C, in the compression zone the temperature was maintained at between 205-215°C and in the melt zone the temperature was maintained at 195-205°C. The die was divided into three zones, the first zone at 185-195°C, the second die zone at 185-195°C and in the final die zone at 195-205°C. The pelletizing head was operated at a setting providing 100 to 300 rpm resulting in a pellet with a diameter of 5 mm and a length as shown in the following Table.

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Table I
PELLETIZER RESULTS

PVC/Wood Fiber	Pelletizer Melt (Temp., °F)	Pellet Length (in.)	Profile Melt (Temp., °F)	Profile Visc.	Sawdust Moisture (%)	Pellet Moisture (%)	Pellet Bulk Density (lb/cc)	Profile Density (g/cc)	Tensile Modulus (psi)	
60/40	0.233	366	2580	4.71	4.83	0.96	0.546 (.006)	1.426	990600	
	60/40	0.233	362	2452	4.71, 4.83	0.96	0.546 (.006)			
	70/30	0.080	375	2274	5.28	1.54	0.454 (.007)	1.43	733300	
	70/30	0.080	376	2299	5.28	1.54	0.454 (.007)	1.435	820100	
	50/50	0.084	382	2327	4.94	1.95	0.347 (.002)	1.367	697600	
	70/30	0.195	385	2431		0.93	0.595 (.005)	1.427	752900	
	70/30	0.195	378	2559		0.93	0.595 (.005)	1.433	787600	
	60/40	0.089	377	1985	5.36	1.33	0.418 (.003)	1.423	1103000	
	60/40	0.089	374	2699	5.36	1.33	0.418 (.003)	1.408	815800	
	50/50	0.201	367	2541	5.33	2.09	0.462 (.004)			
	50/50	0.201	366	2670	5.33	2.09	0.462 (.004)	1.397	724300	
	60/40	0.247	374	1948	4.62	1.03	0.466 (.009)	1.426	860000	
	60/40	0.247	370	2326	4.62	1.03	0.466 (.009)	1.433	996700	
	60/40	0.103	373	1605	5.53	1.57	0.387 (.005)	1.431	985400	
	60/40	0.103	381	2221	5.53	1.57	0.387 (.005)	1.435	855800	
	70/30	0.364	0.202	376	1837	5.25	1.50	0.429 (.010)	1.433	868300
	70/30	0.364	0.202	378	2376	5.25	1.50	0.429 (.010)	1.434	798100
	70/30	0.367	0.085	374	1593		1.48	0.378 (.002)	1.438	714200
	70/30	0.367	0.085	375	2145		1.48	0.378 (.002)	1.439	765000
	50/50	0.67	0.177	371	2393	5.08, 5.51	1.61	0.434 (.007)	1.408	883200
	50/50	0.67	0.177	371	3008	5.08, 5.51	1.61	0.434 (.007)	1.528	1029000
	50/50	366	0.085	370	2666		2.01	0.438 (.003)	1.405	922100
	50/50	366	0.085	369	2257		2.01	0.438 (.003)	1.383	922600

In Table I, the composite material is made from a polyvinyl chloride known as GEON 427 obtained from B.F. Goodrich Company. The wood fibre is sawdust by-product of milling soft woods in the manufacture of wood windows at Andersen Corporation, Bayport, Minnesota. The modulus for natural polyvinyl chloride measured similarly to the composite materials is about 430,000 psi (2900 MPa). The Young's modulus is measured using an Instron Model 450S (Series 9 software) automated materials testing system and uses an ASTM method D-638. Specimens are milled to specifications of the test and are measured at 50% relative humidity, 73°F with

a cross set speed of $0.5 \text{ cm} \cdot \text{min}^{-1}$ ($0.200 \text{ in}/\text{min}$).

Table II contains tensile modulus results from composite materials made using injection moulding technology. The Table shows a substantial increase in tensile modulus when compared to non-composite PVC.

It has been found that the preferred pellet of the invention displays, in a formed structural member, a Young's modulus of at least 500,000 psi (3440 MPa) and commonly falls in the range greater than about 800,000 psi (5500 MPa), preferably between 800,000 and $2.0 \times 10^6 \text{ psi}$ (5500 and 13.8 MPa). Further, the coefficient of thermal expansion of the material is well matched to a compromise between aluminum, PVC and wood products and ranges from about 2.9 to $3.2 \times 10^{-4} \text{ }^\circ\text{C}^{-1}$ (1.6 to $1.8 \times 10^{-5} \text{ in./in.}^\circ\text{F}$). It is believed that the superior properties of the structural members made from the composite or pellet of the invention are in large part due to the nature of the pellet set forth in the Table above. It is considered that the Table clearly shows that the polyvinyl chloride and wood fibre can be combined at various proportions under a variety of temperature conditions to produce a regular pellet. The pellet then can be used in further extrusion processes to form a useful extruded structural member useful in the manufacture of environmentally sensitive windows and doors. The composite is a superior replacement for wood because it has similar mechanical properties but attains a dimensional stability and resistance to rot, and insect damage not attainable by wood products.

Table II

<u>Injection Moulded Samples</u>		
DESCRIPTION	TENSILE MODULUS (ksi)	STANDARD DEVIATION (ksi)
High melt/large pellet/40%	1205	242.4
PVC	488.8	28.4
High melt/small pellet/40%	1232	133.3

Features of polymer/wood composite materials, and components and members made from such materials, are disclosed in US patent applications numbers 07/938604, 07/938364 and 07/938365, and the European patent applications which claim priority from those applications which are being filed with this application. Reference is to be made to the specifications of those applications for information regarding those features.

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Pellet	Small	Large	Small	Large	Small	Large	Small	Large
	1	1	2	2	3	3	2	3
Lg. Pellet	Lg. Pellet	Lg. Pellet	PVC	PVC	Sm. Pellet	Sm. Pellet		
<u>402</u>	<u>402</u>	<u>30.0</u>	<u>30.0</u>	<u>402</u>	<u>402</u>	<u>402</u>	<u>402</u>	<u>402</u>
STD. CYCLE:								
Inj. Past Time	---	---	---	---	---	---	10.0 sec.	30.0 sec.
Inj. Boost Time	2.5	3.0	2.5	3.0	3.0	3.0	---	---
Pill Time	1.0 to	1.2 to	1.2	1.2	1.6	1.6	1.2	1.6
Inj. Hold Time	1.2	1.5	1.2	1.5	1.6	1.6	1.2	1.6
Mold Closed Time	12.0	12.0	12.0	12.0	12.0	12.0	12.0	12.0
Mold Open Time	25.0	25.0	25.0	25.0	25.0	25.0	25.0	25.0
Inj. Fast Pressure	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
Inj. Boost Pressure	1800 Psi	1850 Psi	1850 Psi	1850 Psi	1350 Psi	1350 Psi	1600 Psi	1700 Psi
Inj. Mold Pressure	800 Psi	800 Psi	800 Psi	800 Psi	1000 Psi	1000 Psi	1000 Psi	1000 Psi
Back Pressure	30 lbs.	30 lbs.	30 lbs.	30 lbs.	30 lbs.	30 lbs.	30 lbs.	30 lbs.
Inj. Speed	w/o	w/o	w/o	w/o	w/o	w/o	w/o	w/o
Screw Speed	Fast	Fast	Fast	Fast	Fast	Fast	Slow	Slow
Feed	Slow	Slow	Slow	Slow	Slow	Slow	2"	2"
Cushion	1 1/2"	1 1/2"	1 1/2"	1 1/2"	1 1/2"	1 1/2"	1 1/2"	1 1/2"
Decompress	Off	Off	Off	Off	Off	Off	Off	Off
Front Zone	340°F	340°F	340°F	340°F	340°F	340°F	340°F	340°F
Rear Zone	330	330	330	330	330	330	330	330
Melt Temp.	360 to	360 to	360 to	360 to	360 to	360 to	360 to	360 to
Nozzle Radius	370	370	370	370	370	370	370	370
Nozzle Orifice	Same as Mold	None	None	None	---	---	---	---
Core Seq.	E, J, Stroke	Max.	Max.	Max.	---	---	---	---
K. O. Bar Length	The Backs Std.	---	---	---	---	---	---	---
K. O. Stud Length	5/8"	5/8"	5/8"	5/8"	5/8"	5/8"	5/8"	5/8"
Mold Open Stroke	16"	16"	16"	16"	16"	16"	16"	16"
Multi Stroke	2	2	2	2	2	2	2	2
Die Slow	To Fast - Med.	Tower	Tower	Tower	Tower	Tower	Tower	Tower
A-Side Water								
B-Side Water								
Port Settings								
Full Open	Auto	Auto	Auto	Auto	Auto	Auto	Auto	Auto

Claims

55 1. A composite formed as a pellet, which is formed from a composition which comprises:

- (a) at least about 30 wt % of a polymer comprising vinyl chloride;
- (b) at least about 30 wt % of wood fibre having a minimum thickness of 1 mm and a minimum length of 3 mm and a minimum aspect ratio of about 1.8; and

(c) less than about 8 wt % by weight water;
 in which the wood fibre is dispersed throughout a continuous polymer phase.

2. A polymer and wood fibre composite capable of formation into a structural member, comprising:
 5 (a) at least about 30 wt % of a polymer comprising vinyl chloride;
 (b) at least about 30 wt % of wood fibre having a minimum thickness of 1 mm and a minimum length of 3 mm and an aspect ratio of about 1.8; and
 (c) less than about 8 wt % water;
 in which the wood fibre is dispersed throughout a continuous polymer phase.

10 3. A composite formed from a composition comprises:
 (a) at least about 30 wt % of a polymer comprising vinyl chloride;
 (b) at least about 30 wt % of wood fibre having a minimum width of 0.3 mm and a minimum length of 1 mm and an aspect ratio of greater than about 1.8;
 15 (c) about 0.01 to 25 wt % of an intentionally recycled impurity comprising thermoplastic polymer, adhesive, paint, or thermoplastic resin, or mixtures thereof; and
 (d) less than about 10 wt % water;
 in which the wood fibre is dispersed throughout a continuous polymer phase.

20 4. A composite as claimed in any one of claims 1 to 3, in which the polymer comprises a vinyl chloride homopolymer, preferably with a molecular weight of about $90,000 \pm 50,000$.

5. A composite as claimed in any one of claims 1 to 3, in which the polymer comprises a vinyl chloride co-polymer, preferably with a molecular weight of about $88,000 \pm 10,000$.

25 6. A composite as claimed in any one of claims 1 to 5, in which the wood fibre comprises a by-product of milling or sawing wooden members, preferably sawdust.

7. A composite as claimed in any one of claims 1 to 6, in which the composition additionally comprises a compatibilising agent.

30 8. A composite as claimed in any one of claims 1 to 7, in which the wood fibre has a fibre width of about 0.3 to 1.5 mm.

9. A composite as claimed in any one of claims 1 to 8, in which the wood fibre has a fibre length of about 1 to 10 mm.

35 10. A composite as claimed in any one of claims 1 to 9, in which the wood fibre has an aspect ratio of about 2 to 7.

11. A composite as claimed in any one of claims 1 to 10, in which the water comprises at least about 0.01 wt % of the pellet.

40 12. A composite as claimed in any one of claims 1 to 11, in which the water comprises less than about 5 wt %, preferably less than about 3.5 wt %, of the pellet.

45 13. A composite as claimed in any one of claims 1 to 12, which comprises from about 0.01 to 25 wt % of intentionally recycled impurity, comprising at least one of a thermoplastic polymer, an adhesive, and a thermoplastic resin.

50 14. A composite as claimed in claim 13, which includes one or more of (a) upto about 5 wt % of a hot melt adhesive composition, (b) upto about 5 wt % of a paint, and (c) 0.01 to 15 wt % of recycled polyvinyl chloride cladding which comprises about 9.0 to 99.9 wt % polyvinyl chloride and about 0.01 to 2 wt % of a pigment.

55 15. A composite as claimed in any one of claims 1 to 14, which is a cylindrical extruded pellet having a radius of about 1 to 5 mm and a length of about 1 to 10 mm.

16. A composite as claimed in any one of claims 1 to 14, which is a cylindrical linear extrudate having a radius of about 1 to 5 mm.

17. A composite as claimed in any one of claims 1 to 16, in which the amount of the polymer in the blend is more than about 40%, more preferably more than about 45%.
- 5 18. A composite as claimed in any one of claims 1 to 17, in which the amount of the wood fibre in the blend is more than about 35%.
19. A composite as claimed in any one of claims 1 to 18, in which the amount of polymer in the blend is less than about 70%, preferably less than about 65%.
- 10 20. A composite as claimed in any one of claims 1 to 19, in which the amount of wood fibre in the blend is less than about 50%.

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